### Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims:

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14 15

#### 1 - 30 (cancelled)

- 31 (currently amended): A method for treating an ophthalmic disorder in a 2 mammal, said method comprising administering to the eye of said mammal a lipid formulation, 3 said lipid formulation comprising: 4 a lipid phase, said lipid phase comprising a phospholipid and a modifying agent, 5 wherein said modifying agent is a member selected from the group consisting of cationic 6 lipid[[s]] and mucoadhesive compounds; 7 an aqueous phase; and 8 a therapeutic agent, wherein the therapeutic agent is diclofenac, or a
- 9 pharmaceutically acceptable salt thereof:

10 wherein said therapeutic agent in said lipid formulation is useful for treating said 11 ophthalmic disorder;

wherein said lipid formulation comprises about 0.001 to about 10.000 wt % of said lipid phase and about 90.000 wt % to about 99.999 wt % of said aqueous phase, and wherein said lipid phase comprises 0.1 to 90.0 wt% of the therapeutic agent, 0.01 to 10 wt% 98.8 wt% phospholipid, 0.1 to 10 wt % modifying agent and 0.1 to 10 wt% antioxidant.

- 1 32 (original): The method in accordance with claim 31, wherein said ophthalmic 2 disorder is post-operative pain.
- 1 33 (original): The method in accordance with claim 31, wherein said ophthalmic 2 disorder is ocular inflammation

1	34 (previously presented): The method in accordance with claim 33, wherein
2	said ocular inflammation results from a member selected from the group consisting of iritis,
3	conjunctivitis, seasonal allergic conjunctivitis, acute and chronic endophthalmitis, anterior
4	uveitis, uveitis associated with systemic diseases, posterior segment uveitis, chorioretinitis, pars
5	planitis, ocular lymphoma, pemphigoid, scleritis, keratitis, severe ocular allergy, corneal abrasion
6	and blood-aqueous barrier disruption.
1	35 (original): The method in accordance with claim 31, wherein said ophthalmic
2	disorder is post-operative ocular inflammation.
1	36 (original): The method in accordance with claim 35, wherein said post-
2	operative ocular inflammation results from a member selected from the group consisting of
3	photorefractive keratectomy, cataract removal surgery, intraocular lens implantation and radial
4	keratotomy.
1	37 (original): The method in accordance with claim 31, wherein said ophthalmic
2	disorder is a fungal or bacterial infection.
1	38 (original): The method in accordance with claim 31, wherein said ophthalmic
2	disorder is herpes ophthalmicus.
1	39 (original): The method in accordance with claim 31, wherein said ophthalmic
2	disorder is endophthalmitis.
1	40 (original): The method in accordance with claim 31, wherein said ophthalmic
2	disorder is intraocular pressure.
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	41 - 42 (cancelled)
1	43 (currently amended): A method for treating or preventing ocular
2	inflammation, paracentesis-induced miosis, cystoid macular edema and mydriasis, said method
3	comprising administering a therapeutically effective amount of one or more a first non-steroidal

- 4 anti-inflammatory drug and optionally a second non-steroidal anti-inflammatory drugs
  5 encapsulated or contained within a liposome formulation, said liposome formulation comprising
  6 0.001 to 10.000 wt% lipid phase, and 90.000 to 99.999 wt% aqueous phase, wherein said lipid
  7 phase comprises 0.1 to 90.0 wt% of said first anti-inflammatory drug, 0.01 to 10-wt% 98.8 wt%
  8 phospholipid, 0.1 to 10\_wt%\_modifying agents and 0.1 to 10 wt% antioxidant;
  9 wherein said modifying agent is a cationic lipid;
  10 wherein said first non-steroidal anti-inflammatory drug is diclofenac, or a
- pharmaceutically acceptable salt thereof.

  1 44 (original): The method in accordance with claim 43, wherein said liposome
- 1 44 (original): The method in accordance with claim 43, wherein said liposome 2 formulation is applied topically, resulting in the transcorneal or transscleral passage or 3 introduction of one or more non-steroidal anti-inflammatory drugs into the eye.

# 45 - 46 (cancelled)

1 47 (currently amended): The method in accordance with claim 46, wherein said
2 second non-steroidal anti-inflammatory drugs are selected from the group consisting of
3 ketoprofen, flurbiprofen, ibuprofen, dielofenae, ketorolac, nepafenac, amfenac and suprofen.

## 48 (cancelled)

- 1 49 (previously presented): The method in accordance with claim 43, wherein
  2 said ocular inflammation is a symptom of iritis, conjunctivitis, seasonal allergic conjunctivitis,
  3 post-operative inflammation, acute and chronic endophthalmitis, anterior uveitis, uveitis
  4 associated with systemic diseases, posterior segment uveitis, chorioretinitis, pars planitis, ocular
  5 lymphoma, pemphigoid, scleritis, keratitis, severe ocular allergy, corneal abrasion, blood6 aqueous barrier disruption or ocular trauma.
- 50 (original): The method in accordance with claim 49, wherein said postoperative inflammation is caused by photorefractive keratectomy, cataract removal surgery, intraocular lens implantation or radial keratotomy.

51 - 52 (cancelled)